12-20-04

Attorney's Docket No.: 16863-002001

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Serial No.: 10/622,003

Art Unit: 1644

Examiner: Unknown

Filed : July 16, 2003

Title : PREPARATION OF FULLY HUMAN ANTIBODIES

Mail Stop Amendment Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

In accordance with the duty of disclosure as set forth in 37 C.F.R. §1.56, Applicants hereby submit the following information in conformance with 37 C.F.R. §§ 1.97 and 1.98. Pursuant to 37 C.F.R. § 1.98, a copy of each of the documents cited is enclosed.

Foreign Patent

1. WO 01/00678 A1, published January 4, 2001

Articles

- 2. Dercamp, C., et al. (2002). Depletion of human NK and CD8 cells prior to in vitro H1N1 flu vaccine stimulation increases the number of gamma interferon-secreting cells compared to the initial undepleted population in an ELISPOT assay. Clin Diagn Lab Immunol. 9(2):230-235.
- 3. Kobayashi, Y., et al. (2000). Cell-type specificity of l-leucyl l-leucine methyl ester. Biochem Biophys Res Commun. 272(3):687-690.
- 4. Ohlin, M., et al. (1992). Human MoAbs produced from normal, HIV-1-negative donors and specific for glycoprotein gp120 of the HIV-1 envelope. Clin Exp Immunol. 89(2):290-295.
- 5. Puhlmann, C.M., and Anderer, F.A. (1995). Optimizing production of human monoclonal IgG antibodies by *in vitro*-primed human PBMC: influence of CD56⁺ NK cell depletion. Hybridoma. 14(4):391-396.

CERTIFICATE OF MAILING BY EXPRESS MAIL		
Express Mail Label No	EV584750910US	
	December 17, 2004	
Express Mail Label No		

Date of Deposit

Applicant: Li-Te Chin Attorney's Docket No.: 16863-002001

Serial No. : 10/622,003 Filed : July 16, 2003

Page : 2 of 3

6. Trujillo, J.R., et al. (1998). Shared antigenic epitopes on the V3 loop of HIV-1 gp120 and proteins on activated human T cells. Virology. 246(1):53-62.

7. Zwick, M.B., et al. (2003). A novel human antibody against human immunodeficiency virus type 1 gp120 is V1, V2, and V3 loop dependent and helps delimit the epitope of the broadly neutralizing antibody immunoglobulin G1 b12. J Virol. 77(12):6965-6978.

These documents are being submitted before a first Office Action on the merits; therefore, no fee is required under 37 C.F.R. § 1.97(b). In the event an Office Action is mailed by the United States Patent and Trademark Office prior to receipt of this Supplemental Information Disclosure Statement, Applicants hereby make the statement specified in 37 C.F.R. §1.97(e) that each document contained herein was first cited in the European Search Report for the corresponding European application (EP 04 01 6838.7) within three (3) months of the filing date of this Supplemental Information Disclosure Statement. Therefore, no fee is required under 37 C.F.R. § 1.97(c). A copy of this European Search Report is also enclosed herewith.

By citing the above references, Applicants do not acquiesce or admit that any of these documents is "prior art" under 35 U.S.C. Applicants specifically reserve the right, where appropriate, to antedate any of the cited documents by an appropriate showing under 37 C.F.R. §1.131, §1.604, §1.608 or any other suitable means.

To assist the Examiner, the documents are listed on the attached form PTO-1449. It is respectfully requested that an Examiner initialed copy of this form be returned to the undersigned.

Applicant: Li-Te Chin Attorney's Docket No.: 16863-002001

Serial No. : 10/622,003 Filed : July 16, 2003

Page : 3 of 3

Please apply any charges or credits to Deposit Account No. 06-1050.

Respectfully submitted,

Date: December 17, 2004

Ping F. Hwung Reg. No. 44,164

Fish & Richardson P.C. 500 Arguello Street, Suite 500 Redwood City, California 94063 Telephone: (650) 839-5070 Facsimile: (650) 839-5071

50251945.doc

Substitute Form PTO-1449 (Moduled)

U.S. Department of Commerce Patent and Trademark Office

Attorney's Docket No. 16863-002001 Application No. 10/622,003

Supplemental Information Disclosure Statement **by Applicant** (Use several sheets if necessary)

Li-Te Chin

Filing Date

Applicant

Group Art Unit

(37 CFR §1.98(b))

July 16, 2003

1644

U.S. Patent Documents							
Examiner Initial	Desig. ID AA	Document Number	Publication Date	Patentee	Class	Subclass	Filing Date If Appropriate
	AB						
	AC						

Foreign Patent Documents or Published Foreign Patent Applications								
Examiner	Desig.	Document	Publication	Country or			Trans	slation
Initial	ID	Number	Date	Patent Office	Class	Subclass	Yes	No
	AD	WO 01/00678 A1	01/04/2001	EPO				
	AE							
	AF							

	Other Documents (include Author, Title, Date, and Place of Publication)				
Examiner	Desig.				
Initial	ID	Document			
	AG	Dercamp, C., et al. (2002). Depletion of human NK and CD8 cells prior to in vitro H1N1 flu vaccine stimulation increases the number of gamma interferon-secreting cells compared to the initial undepleted population in an ELISPOT assay. Clin Diagn Lab Immunol. 9(2):230-235.			
	AH	Kobayashi, Y., et al. (2000). Cell-type specificity of l-leucyl l-leucine methyl ester. Biochem Biophys Res Commun. 272(3):687-690.			
	AI	Ohlin, M., et al. (1992). Human MoAbs produced from normal, HIV-1-negative donors and specific for glycoprotein gp120 of the HIV-1 envelope. Clin Exp Immunol. 89(2):290-295.			
	AJ	Puhlmann, C.M., and Anderer, F.A. (1995). Optimizing production of human monoclonal IgG antibodies by in vitro-primed human PBMC: influence of CD56+ NK cell depletion. Hybridoma. 14(4):391-396.			
	AK	Trujillo, J.R., et al. (1998). Shared antigenic epitopes on the V3 loop of HIV-1 gp120 and proteins on activated human T cells. Virology. 246(1):53-62.			
	AL	Zwick, M.B., et al. (2003). A novel human antibody against human immunodeficiency virus type 1 gp120 is V1, V2, and V3 loop dependent and helps delimit the epitope of the broadly neutralizing antibody immunoglobulin G1 b12. J Virol. 77(12):6965-6978.			

Examiner Signature	Date Considered			
	Date Considered			
EXAMINER: Initials citation considered. Draw line through citation if not in conformance and not considered. Include copy of this form with				
novi communication to continue to continue to the continue to				
next communication to applicant.				